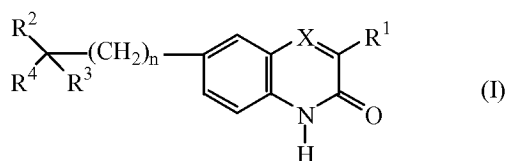


Listing of Claims:

This listing of claims replaces all prior versions, and listings, of claims in the captioned application.

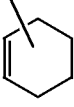
1. (Cancelled)
2. (Currently Amended) A compound of formula (I),



the *N*-oxide forms, the pharmaceutically acceptable addition salts and the stereo-chemically isomeric forms thereof, wherein

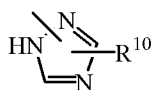
R^1 is C_{1-6} alkyl

R^2 is hydrogen or hydroxy or taken together with R^4 may form $=O$;

R^4 is hydrogen, C_{1-6} alkyl, furanyl, pyridinyl, aryl C_{1-6} alkyl or  ;

n is 0 or 1;

X is N or CR^5 , wherein R^5 is hydrogen;



R^3 is $-(CH_2)_s-$ NR⁶R⁷ or is (c-4) ;

s is 0, 1 or 2;

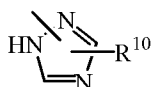
R^6 is $-CHO$, C_{1-6} alkyl, piperidinyl C_{1-6} alkyl, arylcarbonylpiperidinyl C_{1-6} alkyl or aryl C_{1-6} alkyl(C_{1-6} alkyl)amino C_{1-6} alkyl;

R^7 is hydrogen or C_{1-6} alkyl;

and each R^{10} independently is hydrogen, C_{1-6} alkyl or C_{1-6} alkyloxy C_{1-6} alkylamino,

aryl is phenyl or phenyl substituted with halo, C_{1-6} alkyl or C_{1-6} alkyloxy;

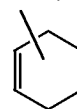
with the proviso that when



n is 0, X is N, R^2 is hydrogen, R^3 is (c-4) and is attached with a nitrogen atom, and R^{10} is hydrogen; then
 R^4 is other than C_{1-6} alkyl or pyridinyl.

3. (Previously Presented) A compound according to claim 2 wherein
 n is 0; X is N or CR^5 , wherein R^5 is hydrogen; R^1 is C_{1-6} alkyl;
 R^2 is hydrogen or hydroxy or taken together with R^4 may form $=O$; R^3 is $-(CH_2)_8-NR^6R^7$;

s is 0 or 1; R^6 is $-CHO$ or C_{1-6} alkyl; and R^4 is hydrogen, C_{1-6} alkyl or



4. (Previously Presented) A compound selected from the group consisting of:

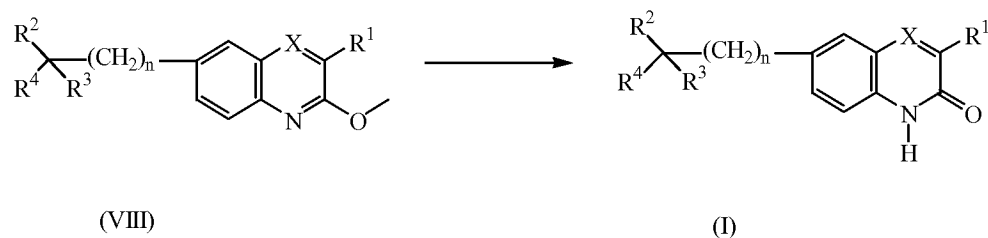
<p>compound 1</p>	<p>compound 5</p>
<p>compound 7</p>	<p>compound 3</p>
<p>compound 17</p>	

and the *N*-oxide forms, the pharmaceutically acceptable addition salts and the stereochemically isomeric forms thereof.

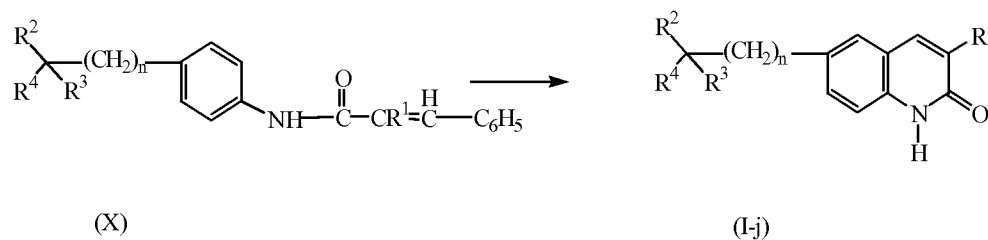
5. (Cancelled)
6. (Previously Presented) A pharmaceutical composition comprising pharmaceutically acceptable carriers and as an active ingredient a therapeutically effective amount of a compound as claimed in claim 2.
7. -11. (Cancelled).

12. (Currently Amended) A combination of a compound with a cancer treating chemotherapeutic agent wherein said compound is a compound of formula (I) according to Claim 2.

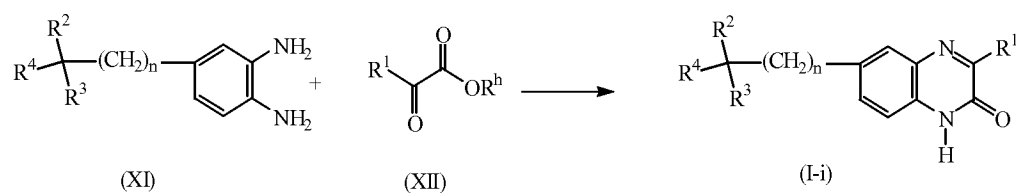
13. (Previously Presented) A process for preparing a compound as claimed in claim 2, comprising: a) hydrolysis of intermediates of formula (VIII),



b) cyclization of intermediates of formula (X),



or c) condensation of an ortho-benzenediamine of formula (XI) with an ester of formula (XII) wherein R^h is C₁₋₆alkyl, into compounds of formula (I), wherein X is N, herein referred to as compounds of formula (I-i),



14. (Cancelled)

15. (Previously Presented) A pharmaceutical composition comprising pharmaceutically acceptable carriers and as an active ingredient a therapeutically effective amount of a compound as claimed in claim 3.

16. (Previously Presented) A pharmaceutical composition comprising pharmaceutically acceptable carriers and as an active ingredient a therapeutically effective amount of a compound as claimed in claim 4.

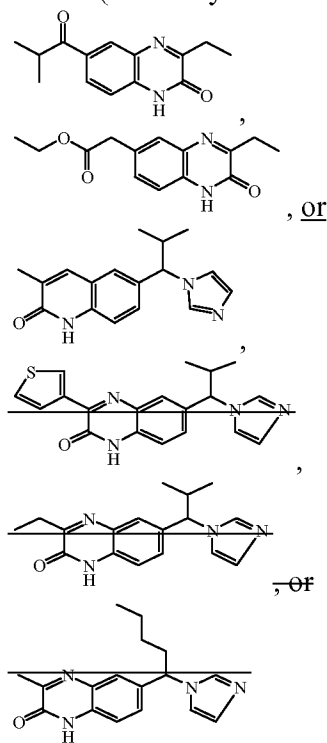
17. - 26. (Cancelled)

27. (Currently Amended) A combination of a compound with a cancer treating chemotherapeutic agent wherein said compound is a compound of claim 3.

28. (Currently Amended) A combination of a compound with a cancer treating chemotherapeutic agent wherein said compound is a compound of claim 4.

29. - 30. (Cancelled)

31. (Currently Amended) A compound selected from



and the *N*-oxide forms and the pharmaceutically acceptable addition salts thereof.

32. (Previously Presented) A pharmaceutical composition comprising pharmaceutically acceptable carriers and as an active ingredient a therapeutically effective amount of a compound as claimed in claim 31.

33. (Previously Presented) A combination of a compound with a chemotherapeutic agent wherein said compound is a compound of claim 31.